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Terms	Documents
6200577.pn.	2

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<u>L10</u>	6200577.pn.	2	<u>L10</u>
<u>L9</u>	l7 and lipid\$	0	<u>L9</u>
<u>L8</u>	L7 and aggregate\$	0	<u>L8</u>
<u>L7</u>	6086902.pn.	2	<u>L7</u>
<u>L6</u>	l1 not l5	45	<u>L6</u>
<u>L5</u>	l1 same lipid\$	82	<u>L5</u>
<u>L4</u>	aggregate\$ same (nucleic adj acid\$) same vp22	1	<u>L4</u>
<u>L3</u>	L1 and vp22	6	<u>L3</u>
<u>L2</u>	L1 same vp22	1	<u>L2</u>
<u>L1</u>	aggregate\$ same (nucleic adj acid\$) same deliver\$	127	<u>L1</u>

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Terms	Documents
l1 not l5	45

Database:

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<u>L6</u>	l1 not l5	45	<u>L6</u>
<u>L5</u>	l1 same lipid\$	82	<u>L5</u>
<u>L4</u>	aggregate\$ same (nucleic adj acid\$) same vp22	1	<u>L4</u>
<u>L3</u>	L1 and vp22	6	<u>L3</u>
<u>L2</u>	L1 same vp22	1	<u>L2</u>
<u>L1</u>	aggregate\$ same (nucleic adj acid\$) same deliver\$	127	<u>L1</u>

END OF SEARCH HISTORY

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(c) 2003 Inst for Sci Info

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File 370:Science 1996-1999/Jul W3

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File 399:CA SEARCH(R) 1967-2003/UD=13822

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Set	Items	Description
---	-----	-----
?s aggregate?	(s)	(nucleic acid)
	397734	AGGREGATE?
	74893	NUCLEIC ACID
S1	0	AGGREGATE? (S) (NUCLEIC ACID)
?s vp22	(s)	aggregate?
	846	VP22
	397734	AGGREGATE?
S2	8	VP22 (S) AGGREGATE?
?rd		
...completed	examining	records
S3	5	RD (unique items)
?show files;	ds;t/3,k/all	
File 5:	Biosis Previews(R)	1969-2003/May W4
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Set	Items	Description
S1	0	AGGREGATE? (S) (NUCLEIC ACID)
S2	8	VP22 (S) AGGREGATE?
S3	5	RD (unique items)

>>>KWIC option is not available in file(s): 399

3/3,K/1 (Item 1 from file 5)
DIALOG(R)File 5:Biosis Previews(R)
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06956495 BIOSIS NO.: 000089078502

**THREE-DIMENSIONAL STRUCTURES OF MATURABLE AND ABORTIVE CAPSIDS OF EQUINE
HERPESVIRUS 1 FROM CRYOELECTRON MICROSCOPY**

AUTHOR: BAKER T S; NEWCOMB W W; BOOY F P; BROWN J C; STEVEN A C
AUTHOR ADDRESS: LAB. PHYSIOL. BIOL., NATL. INST. ARTHRITIS MUSCULOSKELETAL
SKIN DIS., BUILDING 6, ROOM 114, BETHESDA, MD. 20892.
JOURNAL: J VIROL 64 (2). 1990. 563-573. 1990
FULL JOURNAL NAME: Journal of Virology
CODEN: JOVIA
RECORD TYPE: Abstract
LANGUAGE: ENGLISH

...ABSTRACT: which is smeared out in the reconstruction, implying that its distribution is not icosahedrally symmetric or necessarily consistent from particle to particle.sbd.consists of *aggregates* of *VP22* (46 kDa). From several lines of evidence, we conclude that this protein is located entirely within the capsid shell. These *aggregates* may be the remnants of morphogenetic cores retained in capsids interrupted in the process of DNA packaging.

3/3,K/2 (Item 1 from file: 399)
DIALOG(R)File 399:CA SEARCH(R)
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136241644 CA: 136(16)241644v PATENT
VP22 protein/nucleic acid aggregates for delivery to cells
INVENTOR(AUTHOR): O'Hare, Peter Francis Joseph; Brewis, Neil Douglas;
Normand, Nadia Michelle; Sunassee, Kavitha Renga
LOCATION: UK,
ASSIGNEE: Phogen Limited
PATENT: PCT International ; WO 200220060 A1 DATE: 20020314
APPLICATION: WO 2001GB4057 (20010910) *GB 200022101 (20000908)
PAGES: 31 pp. CODEN: PIXXD2 LANGUAGE: English CLASS: A61K-048/00A;
C12N-015/87B DESIGNATED COUNTRIES: AE; AG; AL; AM; AT; AU; AZ; BA; BB; BG;
BR; BY; BZ; CA; CH; CN; CO; CR; CU; CZ; DE; DK; DM; DZ; EC; EE; ES; FI; GB;
GD; GE; GH; GM; HR; HU; ID; IL; IN; IS; JP; KE; KG; KP; KR; KZ; LC; LK; LR;
LS; LT; LU; LV; MA; MD; MG; MK; MN; MW; MX; MZ; NO; NZ; PH; PL; PT; RO; RU;
SD; SE; SG; SI; SK; SL; TJ; TM; TR; TT; TZ; UA; UG; US; UZ; VN; YU; ZA; ZW;
AM; AZ; BY; KG; KZ; MD; RU; TJ; TM DESIGNATED REGIONAL: GH; GM; KE; LS; MW
; MZ; SD; SL; SZ; TZ; UG; ZW; AT; BE; CH; CY; DE; DK; ES; FI; FR; GB; GR;
IE; IT; LU; MC; NL; PT; SE; TR; BF; BJ; CF; CG; CI; CM; GA; GN; GQ; GW; ML;
MR; NE; SN; TD; TG

3/3,K/3 (Item 2 from file: 399)
DIALOG(R)File 399:CA SEARCH(R)
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135087161 CA: 135(7)87161f PATENT
**Uses of transport-active proteins and fusion polypeptides, especially
with VP22 function, for controlling cell cycle and inhibiting cell
proliferation**
INVENTOR(AUTHOR): O'Hare, Peter Francis Joseph; Normand, Nadia Michelle;
Brewis, Neil Douglas; Phelan, Anne
LOCATION: UK,
ASSIGNEE: Phogen Limited
PATENT: PCT International ; WO 200147960 A1 DATE: 20010705
APPLICATION: WO 2000GB4965 (20001221) *GB 9930519 (19991224)
PAGES: 23 pp. CODEN: PIXXD2 LANGUAGE: English CLASS: C07K-014/035A;
A61K-047/48B; A61P-035/00B DESIGNATED COUNTRIES: AE; AG; AL; AM; AT; AU;
AZ; BA; BB; BG; BR; BY; BZ; CA; CH; CN; CR; CU; CZ; DE; DK; DM; DZ; EE; ES;
FI; GB; GD; GE; GH; GM; HR; HU; ID; IL; IN; IS; JP; KE; KG; KP; KR; KZ; LC;

LK; LR; LS; LT; LU; LV; MA; MB; MG; MK; MN; MW; MX; MZ; NO; NZ; PL; PT; RO; RU; SD; SE; SG; SI; SK; SL; TJ; TM; TR; TT; TZ; UA; UG; US; UZ; VN; YU; ZA; ZW; AM; AZ; BY; KG; KZ; MD; RU; TJ; TM DESIGNATED REGIONAL: GH; GM; KE; LS; MW; MZ; SD; SL; SZ; TZ; UG; ZW; AT; BE; CH; CY; DE; DK; ES; FI; FR; GB; GR; IE; IT; LU; MC; NL; PT; SE; TR; BF; BJ; CF; CG; CI; CM; GA; GN; GW; ML; MR; NE; SN; TD; TG

3/3,K/4 (Item 1 from file: 357)
DIALOG(R)File 357:Derwent Biotech Res.
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0290584 DBR Accession No.: 2002-12431 PATENT

Use of *aggregates* comprising *VP22* protein/polypeptide with the transport function of *VP22* and oligonucleotides/polynucleotides with disaggregating agent, useful for treating or preventing cell proliferation - vector-mediated gene transfer and expression in host cell for recombinant protein production and gene therapy

AUTHOR: O'HARE P F J; BREWIS N D; NORMAND N M; SUNASSEE K R

PATENT ASSIGNEE: PHOGEN LTD 2002

PATENT NUMBER: WO 200220060 PATENT DATE: 20020314 WPI ACCESSION NO.: 2002-304326 (200234)

PRIORITY APPLIC. NO.: GB 200022101 APPLIC. DATE: 20000908

NATIONAL APPLIC. NO.: WO 2001GB4057 APPLIC. DATE: 20010910

LANGUAGE: English

Use of *aggregates* comprising *VP22* protein/polypeptide with the transport function of *VP22* and oligonucleotides/polynucleotides with disaggregating agent, useful for treating or preventing cell proliferation - vector-mediated gene transfer and expression in host cell for recombinant protein...

ABSTRACT: DERWENT ABSTRACT: NOVELTY - Use of *aggregates* comprising *VP22* protein (or a polypeptide with the transport function of *VP22*), and oligonucleotides or polynucleotides with a disaggregating agent (simultaneously or sequentially) to treat target cells by delivering molecules to the cells and/or preventing cell...

... treating target cells to deliver molecules to the cells and/or prevent their proliferation and/or kill them comprising: (a) exposing the cells to the *aggregate* composition cited above; and (b) exposing the cells to the disaggregating agent cited above, which can promote disaggregation of the *aggregate* composition in cells, where steps (a) and (b) are carried out simultaneously or sequentially; (2) a product comprising the *aggregate* composition and the disaggregating agent, as combined preparation for administration of these components, either sequentially or together; (3) a pharmaceutical comprising the *aggregate* composition and the disaggregating agent, in combination with a pharmaceutical excipient; and (4) a cell preparation obtainable by treating the target cells in vitro as cited in the method above.

BIOTECHNOLOGY - Preferred Composition: The *VP22* protein or the polypeptide with the transport function of *VP22* is a fusion protein, which also comprises a non-*VP22* polypeptide sequence. The *VP22* protein or the polypeptide with the transport function of *VP22* is a fusion protein is chemically cross linked to a non-*VP22* molecule. The oligonucleotide or polynucleotide comprises a circular plasmid, and is linked to an additional molecule. The disaggregating agent is a photoactivator and can promote disaggregation of the *aggregate* compositions following illumination with actinic light. Preferably, the disaggregating agent is a phthalocyanine-containing chromophore. The disaggregating agent is also an agent that can promote...

...are cells in vitro or in vivo. The method further comprises exposing the target cells to actinic light after delivery of the disaggregating agent. The *aggregates* and the disaggregating agent are administered separately to target cells in vivo at the same loci or at closely neighboring loci. The *aggregates* and disaggregating agent may also be administered together as a combined preparation to target cells in

vivo. ACTIVITY - Cytostatic; antipsoriatic; dermatological. Fifty microliters of the *aggregate* solution was directly injected into CT26 tumors in mice, and then the mice were observed for signs of distress.

Twenty four hours after administration of the *aggregates*, the mice were anesthetized and half of the CT26 tumors were illuminated for 10 minutes using a cold light source KL2500 LCD. Twenty four hours after illumination, the tumors were removed from the mice. Controls included mice injected with phosphate buffered saline (PBS), MH3 peptide, or *aggregates* formed using *VP22* having greater than 159-301 protein and intercellular adhesion molecule (ICAM) oligonucleotides instead of the *aggregates* described above. Test *aggregates* were found to induce significant apoptosis of the CT26 tumor cells in comparison to controls following excision of the tumors from the mice. MECHANISM OF ACTION - Gene therapy; protein therapy. USE - The *aggregate* composition and disaggregating agent are useful in the manufacture of a medicament for treating diseases or target cells, and/or preventing cell proliferation and/or...

3/3,K/5 (Item 2 from file: 357)

DIALOG(R)File 357:Derwent Biotech Res.

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0260679 DBR Accession No.: 2001-00255 PATENT

***Aggregated* composition suitable for phototherapy or prophylaxis of psoriasis, eczema or skin cancer and for delivering nucleic acids and proteins into cells, comprises transport protein *VP22* and an oligonucleotide - plasmid-mediated gene transfer, expression in Escherichia coli, antisense oligonucleotide, ribozyme and recombinant protein aggregate for cancer therapy**

AUTHOR: O'Hare P F J; Normand N M

CORPORATE SOURCE: Cambridge, UK.

PATENT ASSIGNEE: Phogen 2000

PATENT NUMBER: WO 200053722 PATENT DATE: 20000914 WPI ACCESSION NO.: 2000-594314 (2056)

PRIORITY APPLIC. NO.: GB 9930499 APPLIC. DATE: 19991224

NATIONAL APPLIC. NO.: WO 2000GB897 APPLIC. DATE: 20000310

LANGUAGE: English

***Aggregated* composition suitable for phototherapy or prophylaxis of psoriasis, eczema or skin cancer and for delivering nucleic acids and proteins into cells, comprises transport protein *VP22* and an oligonucleotide**

ABSTRACT: An *aggregated* composition (I) containing a protein having the transport function of a transport protein *VP22*, and an oligonucleotide (antisense or ribozyme) or DNA, is claimed. Also claimed are: making (I) by: mixing a protein with the transport function of *VP22* with the oligonucleotide or DNA; and allowing the obtained mixture to form *aggregates* of 0.1-5 microns; and a cell preparation which has been treated with (I). (I) is useful for preparing a medicament for therapy and for delivering molecules to cells in vitro. The medicament is suitable for phototherapy. The *aggregates* are delivered to target cells such as tumor cells in vivo using e.g. liposome and are useful for treating psoriasis, eczema or skin cancer. In an example, a protein designated 159-301 protein consisting of amino acids 159-301 of the herpes-simplex virus-2 *VP22* protein along with a his6 tag at the C-terminus and a 20-mer phosphorothioate oligonucleotide labeled at the 3'-end with fluorescein was prepared. 159-301 was made in an Escherichia coli expression system expressing a plasmid encoding 159-301 protein. The *aggregates* were then added to HeLa cells and incubated for 12 hr at 37 deg. (28pp)

DESCRIPTORS: plasmid-mediated antisense oligonucleotide, ribozyme, herpes-simplex virus recombinant *VP22* protein *aggregate* gene transfer, expression in Escherichia coli, liposome, fluorescein, appl. psoriasis, eczema, skin cancer gene therapy RNA enzyme herpes virus bacterium lipofection transfection fluorescence tumor DNA...